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(54) Title: **COLLECTOR INTENDED TO BE USED IN DEINKING OF WASTE PAPER ACCORDING TO THE FLOTATION PROCESS**

(57) Abstract

A collector that is intended to be used at deinking of waste paper according to the flotation process. The collector includes a polyester obtained through reaction between polyalkylene glycol, di and/or tricarboxylic acid and/or anhydrides thereof as well as a saturated fatty acid with 12-18 carbon atoms. The polyester shows a molecular weight between 3000-10000.

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COLLECTOR INTENDED TO BE USED IN DEINKING OF WASTE PAPER
ACCORDING TO THE FLOTATION PROCESS

BACKGROUND OF THE INVENTION

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Around the world the opinion for use of the recycled fibre is now growing. A number of countries have already legislated that a certain part of the total paper production shall be constituted of recycled fibre. The methods, which 10 are used today at deinking are the flotation process and the washing process, where the later is mainly applied in North America. The flotation process, which is most common in Europe, technically originates from the mining industry and is the process being most interesting for our object.

15

The deinking can be divided in three different steps; 1) dissolution of recycled fibre and discharging printing-ink from the fibres, 2) dispersing printing-ink in the water phase and 3) separation of the printing-ink. These three 20 steps are included in both flotation process and the washing process. The method, which one uses industrial depends on the requirements and conditions which are found where the activity is located. The object of the flotation process concerning the deinking is to remove the very most amount of 25 printing-ink from a recycled fibre suspension. The first step is to solve the fibre and disengage the printing-ink. When this is accomplished the particles must be modified so that they obtain suitable physical and chemical characteristics so that they afterwards can be lifted to the 30 surface and removed with help of air-bubbles.

Some parameters are important to be consider at the flotation, such as the size of the particle, airflow in the flotation cell, the size of the air-bubbles, temperature, 35 flotation time, the concentration of the recycled fibres in the flotation cell, pH, the concentration of the chemicals as well as the degree of the hardness of the water.

The flotation process is carried out in such a manner that the recycled raw material of the fibre is poured out through mechanical treatment in combination with a concentration of the chemicals. Different methods concerning chemicals/the 5 concentration of the chemicals and method to pour the raw material exist. The purpose of the mechanical treatment is that the fibres should be rubbed against each other and that one in that way, together with the chemicals, should get an efficient discharging of print-ink. However, a too strong 10 mechanical influence causes great damages on the fibres and this is not desirable at the paper-making.

Then comes filtering, whereby the particle shaped impurities are separated, followed of a purification step, generally by 15 means of a vortex cleaner, where very little and heavy particles are separated. In a dispersing step the discharge of printing-ink continues through mechanical treatment and chemical influence. Printing-ink and remaining small particles are separated subsequently in the flotation step. 20 Different flotation technics exist and in the more contemporary ones, one can float all small particles by pressurizing the flotation cell. Small impurities are separated through washing and to increase the brightness of the pulp the bleaching is performed through hydrogen 25 peroxide or hydro sulphite.

Following chemicals are used at the deinking:

Sodium hydroxide. (NaOH)

30 Sodium hydroxide is used to adjust pH to the alkaline level and to saponificate and/or hydrolyse the resin in the printing-ink. The pH at, which the deinking occurs, 9.5 - 11.0, causes that the fibres swell and thereby become more flexible. Adding NaOH to the mechanical pulps makes the 35 pulp to turn yellow and become dark. The concentration of NaOH must be optimized. The alkalinity must be high enough, as mentioned earlier, to obtain good saponification and

hydrolysis of resins, which are included in the printing-inks. A sufficient high pH also gives a good fibre flexibility as well as an optimal function of the hydrogen peroxide. A quandary, as mentioned earlier, is that in the 5 volumes with high lignin content the high alkali content give rise to chromophore groups. These chromophore groups are the reason for the pulp becoming yellow dyed. Experiments done show that the brightness increases with increasing pH when it is increased from 8.6 to 10.2, 10 subsequently the brightness diminishes. Increasing number of magazine paper in the pulp requires additional content of sodium hydroxide. Before the pulp is used for paper making pH is reduced to about 5.5. It is shown that the brightness increases linear with decreasing pH between 8.5 and 5.5. The 15 reason for the brightness increase is not determined entirely but one supposes that it depends on two main factors: agglomeration of the colloidal printing-ink as well as reduction of the number of chromophors.

20 **Hydrogen peroxide. (H_2O_2)**

Hydrogen peroxide is used to discolour the chromophors, which are originate from alkaline pH in the pulp. The peroxide reacts with the sodium hydroxide according to the following:

25



when pH=10.0-11.5 and temperature=40-80°C

30 The perhydroxyl anion (HOO^-) is the active bleach chemical. The concentration of this, is influenced by the following parameters: pH, temperature, dose of hydrogen peroxide and the concentration of the certain metal ions such as Fe, Mn, Cu and others. The influence of the metal ions can be reduced by adding chelate and sodium silicate. Certain 35 microorganisms can also have negative influence by producing peroxide degrading enzymes.

Sodium silicate

Sodium silicate, or water-glass, is a flotation silicate, which consists of SiO_2 and Na_2O in varying mole ratio. Sodium metal silicate has a mole ratio ($\text{Na}_2\text{O}/\text{SiO}_2$) of approximately 5 one and is the one mainly being used within the deinking. This is due to the higher alkalinity of the silicates with the low mole ratio. Actually, the Silicate component is a mixture of many complex polymer silicate anions. By forming colloids with heavy metal ions the sodium silicate prevents 10 a degradation of the hydrogen peroxide. The silicate even prevents the loose printing-ink from reprecipitating on the fibres. The silicate has also a buffering effect on pH. Up to a certain level the brightness of the pulp increases with 15 added concentration of the sodium silicate. However, the concentration of the silicate should be kept as low as possible, since problems with the deinkings in the subsequent equipment have been noted.

Chemicals forming agglomerate

20

The printing-ink from the office waste form thin flake at the deinking. These are to big to be able to be removed by flotation or washing and to little to be able to be removed through filtration. If a chemical that changes the surface 25 structure of the platform is added, they can be made to form agglomerate, which can be removed, e.g. through centrifugation.

The surface-active substances

30

The group of surface-active substances include many subgroups. For instance dispersing substances, collectors, substances with wetting characteristics, substances that prevents redeposition and so on, can be mentioned. As a 35 matter of principle, the surface-active substances being used for deinking will have two important components. A portion of the molecule must be hydrophilic and the other

one must be of hydrophobic quality. The structure of the surface-active substances can be a straight chain, branched chain, they can have charged groups, long or short chain, double or simple bindings and so on.

5

Complex builder

DTPA (diethylentriaminopentaacetic acid) is the most used complex builder but EDTA (ethylendiamintetraacetic acid) is 10 also used. The complex builders act by building solvable complexes of heavy metal ions. The complex prevents these ions from breaking down the hydrogen peroxide. DTPA, which has a structure with five "legs" is a more efficient complex builder than EDTA, which has a structure with four "legs".

15

Today, the waste paper manufacturers largely use a raw material mixture, which normally 30-40% consists of magazine paper and to 60-70% of newspaper. It is possible by sorting magazine paper and newspaper by a reasonably cost-effective 20 method. By knowing, which print methods and printing-inks being used, we can easily understand, which mechanisms and functions that are required in the deinking works. For further operation one must also consider following facts about the printing-inks and printing methods:

- 25 * Pigment consists of very fine carbon particles.
- * The carriers consist of mineral or vegetable oils or as well as dissolver, which evaporates.
- * The printing-ink at printing on newspapers is partly absorbed by the paper and thereby contains oils, which make 30 the colour to not dry completely. The printing-ink on the magazine paper consists of endured pigment on the paper surface, which largely consists of filler. This causes that a smaller amount of colour be adsorbed on the cellulose fibre.
- 35 * Both foregoing types of printing-inks have a pronounced hydrophobic quality.

Earlier, the chemical system at flotation has been described to give an insight into what effect the different added ingredients have on the deinking. When the collector is added, the printing-ink is dispersed in the water phase. Now 5 it is required to modify the particles so that they obtain characteristics, which makes it easy for them to be flotated.

If one makes the small hydrophilic particles more 10 hydrophobic, this gives a certain agglomeration. This yields heavier particles with a hydrophobic surface structure. An important condition for the particle to come in contact with the air-bubble is that the particle has an appropriate size.

15 If the particle is too light, it will follow the flow round the bubble and no effect is obtained. If the particle is too heavy a collision with the air-bubble is obtained, however, the adhesive powers are too weak in proportion to its pulp. Therefore, the particle falls off and the flotation effect 20 does not occur. It is therefore very important that the particle is sufficiently big. Earlier experiments have shown that an optimal size for the particle is about 100 μ m.

25 The hydrophobic surface structure favours an adsorption to air-bubbles. The adsorption can be explained using surface chemistry, whereby through an interface between the air and the water the hydrophobic matters are oriented against the air phase. This occurs because it is advantageous from an 30 energy point of view. To be able to achieve a hydrophobing of the particles, one can add a surface-active chemical.

The conventional type of flotation chemical has required a high degree of hardness of the water to obtain intended effect. This acquires problem with depositions in form of 35 CaCO_3 in the equipment. For a period there has been different alternatives to the original flotation chemicals, for instance so-called synthetic collector, which do not

require a high water hardness.

In SE-B-464 639 a synthetic collector, which consists of a polyester based on a polyalkyline glycol and a di or 5 tricarboxylic acid is described. Tall oil is used of the polymerization reaction to control the chain length of the resulting polymer. The collector produced in this manner is efficient and also non-polluting, as it is biologically degradable. However, there is still place for enhancements, 10 for example the possibility of controlling the polymerization reaction and the chain length of the polyester.

THE OBJECT AND THE MOST IMPORTANT CHARACTERISTICS

15 The object of the invention is to obtain an efficient synthetic collector, where the polymerization reaction can be controlled in a better way to thereby increase the reproducibility and give a more well-defined product. 20 Naturally, the environmental aspects applying the biological degradability should be satisfied.

This, according to the invention has been achieved by including a polyester obtained through reaction between 1) 25 poly alkyline glycol, 2) di- and/or tricarboxylic acid and/- or anhydrides thereof as well as 3) a saturated fatty acid with 12-18 carbon atoms, and, which polyester shows a molecular weight between 3000 - 10000.

30 DESCRIPTION OF THE DRAWINGS

Fig. 1 is a block diagram showing the brightness increase at flotation experiment with a collector according to the invention by different dosages.

35 Fig. 2 is a block diagram showing the brightness increase at flotation experiment with a collector according to the invention and a commercial collector.

DESCRIPTION OF THE INVENTION

The collector is a polyester made of a polyalkylene glycol and a di and/or tricarboxylic acid or its anhydrides. Preferably, as polyalkylene glycol polyethylene glycol, PEG, HO-(CH₂-CH₂O)_n-H and/or polypropylene glycol, PPG, HO-(CH₂-CH₂-CH₂O)_n-H are used. These can occur in several different molecular weights. We have used PEG as average molecular weight 400.

10 Preferably, as di or tricarboxylic acid is used maleic acid, fumaric acid, adipic acid, citric acid, oxalic acid or sebamic acid or its anhydrides. Mixtures can also be used.

15 To obtain a certain chain length a blockage acid at the synthesis is added and in the experiments, already described tall oil (Bevacid-20), myristic acid, lauric acid and stearic acid has been used. Through the polycondensation between PEG and maleic acid a chain with a relative 20 hydrophilic structure caused by carboxyl groups and oxygen atoms is obtained. It is also desired a strong hydrophobic part on the molecule and this characteristic is obtained by choosing a hydrophobic blockage acid. When one varies the polymerisation ratio and the length of the blockage acid, 25 different efficiency with the collector is obtained.

Tall oil consists of 25-30% resin acids and 70-75% fatty acids. The portion of fatty acid in the tall oil is approximately 5% saturated, approximately 25% consists of 30 oil acid and approximately 70% consists of linolenic acid. The mixture of different fatty acids makes it difficult to decide the appearance of the collector, but if estimation is done one can establish that the half the tall oil consists of linolenic acid.

35 According to the invention three saturated fatty acids with 18, 14 and 12 carbon atoms respectively, i.e. stearic acid,

myristic acid and lauric acid have been tested as blockage acids. To be able to study how the chain length of the molecules affected the collector function, synthesis with calculated molecular weights (Mn) of 3000, 5000 and 7000 5 were made. The average molecular weight should be between 3000-10000.

A polyester was formed by gradually polymerization between two bi or multifunctional monomers. By a bifunctional 10 monomer is meant a molecule with two functional groups. The reaction is also called condensation polymerization because the water is separated.

Example:

15



Generally, condensation polymers are formed through gradually intermolecular condensation of reactive groups. 20 Condensation polymers can also be formed through additional polymerisation. Bifunctional monomers give linear polymers, while polyfunctional gives branched or cross linked polymers. To interrupt the reaction one usually uses a monofunctional acid, which is called blockage acid.

25

To obtain a linear polymeric with high molecular weight the exchange must be much higher. A ratio of a transformer of 90%, for instance gives only one average of 10 repeating units in the chain. High exchange is obtained for instance 30 by means of displacing the equilibrium in the above reaction to right. It is also important that the monomers are added in equimolar quantities and there are no impurities present. The temperature should not be too high, since there is a risk of bireaction, which leads to cross-linkage and 35 branching. To obtain a faster polymerization one usually adds a strong acid as catalyst, for instance sulphuric acid or p-toluensulphonic acid. It is also possible to use

certain tin organic combinations as catalysts.

The prerequisite for gradually polymerization is following:

- 5 - any of the two present reactants can react.
- the concentration of the monomer decreases fats in the beginning of the reaction.
- the molecular weight of the polymer increases whole the time during the reaction.
- 10 - long reaction time is necessary to obtain high molecular weight.

Synthesis of the collector

15 The included raw materials were poured in a flange flask, the heating was started and the catalyst added. Timekeeping was started when the catalyst was added. Nitrogen was conducted in the mixture to obtain a certain stirring as well as to prevent oxygen supply. To improve the stirring a magnetic agitator was used. Water suction was used since low 20 pressure lowers both partial pressures of water vapour in the flask and the boiling point at the reaction mixture.

25 To be able to obtain a controlled temperature increase as well as be able to carry out the synthesis at a constant temperature an adjustable transformer was connected to the heat casing. The synthesis went on in about three hours. Sometimes the synthesis was interrupted earlier because the mixture became too viscous or a certain acid ratio was obtained. At regular intervals samples for control of the 30 acid ratio were taken.

35 The synthesis was accomplished at 150, 175 and 200°C respectively to be able to study the impact of the synthesis temperature on the molecular weight. For each synthesis a double sample is performed to control the reproducibility for the synthesis. After each synthesis the amount of the given water was measured and this condensate was

controlled by means of FTIR. The synthesis product was characterized by means of FTIR (Fourier Transform Infrared Spectroscopy), acid ratio, viscosity and in a certain case even by means of NMR (Nuclear Magnetic Resonance), clouding 5 point and PLC (High Pressure Liquid Chromatography).

When a surface-active substance is present as a diluted water solution, during the clouding point, the molecules are dissolved well. If the temperature is increased, the 10 molecules begin to approach each other more and more. Finally it is visible when the solution becomes milky. This is called the clouding point. Because of this it is important that the molecule is designed with regard to the flotation temperature. A number of different collectors were 15 produced according to following recipe and reaction conditions:

	<u>COLLECTOR 1, 2, 4</u>	<u>COLLECTOR 3, 5</u>	<u>COLLECTOR 6, 7</u>
20	Reaction time: 240, 210, 210 min resp. 150°C Maleine acid 102 g PEG 400 461 g Tall oil 35 g	Reaction time: 165 and 150 min resp. 175°C otherwise as 1, 2	Reaction time: 180 and 150 min 200°C otherwise as 1, 2
25	<u>COLLECTOR 8, 9</u> Reaction time: 210 and 195 min rasp. 150°C Malein acid 116 g PEG 400 400 g Tall oil 40 g	<u>COLLECTOR 10, 11</u> Reaction time: 155 and 180 min rasp. 175°C otherwise as 8, 9	<u>COLLECTOR 12, 13</u> Reaction time: 188 and 180 min 200°C otherwise as 8, 9
30			
35	<u>COLLECTOR 14</u> Reaction time: 180 min. 175°C Maleine acid 102 g PEG 400 461 g Stearic acid 40 g	<u>COLLECTOR 15, 17</u> Reaction time: 180 min. 150°C otherwise as 14	<u>COLLECTOR 16</u> Reaction time: 150 min. 200°C otherwise as 14
40			

	<u>COLLECTOR 18</u>	<u>COLLECTOR 19</u>	<u>COLLECTOR 20</u>
	Reaction time: 210 min. 150°C Maleine acid 116 g PEG 400 400 g Tall oil 103 g	Reaction time: 180 min. 150°C Maleine acid 116 g PEG 400 400 g Stearic acid 108 g	Reaction time: 210 min. 150°C Maleine acid 116 g PEG 400 400 g Tall oil 57 g
5			
10	<u>COLLECTOR 21</u> Reaction time: 180 min. 150°C Maleine acid 116 g PEG 400 400 g Stearic acid 60 g	<u>COLLECTOR 22</u> Reaction time: 180 min. 150°C Maleine acid 116 g PEG 400 400 g Myristic acid 34 g	<u>COLLECTOR 23</u> Reaction time: 180 min. 150°C Maleine acid 116 g PEG 400 400 g Lauric acid 30 g
15			
20	<u>COLLECTOR 24</u> Reaction time: 180 min. 150°C Maleine acid 116 g PEG 400 400 g Lauric acid 76 g	<u>COLLECTOR 25</u> Reaction time: 240 min. 150°C Fumaric acid 116 g PEG 400 400 g Myristic acid 34 g	<u>COLLECTOR 26</u> Reaction time: 150 min. 200°C Fumaric acid 102 g PEG 400 461 g Stearic acid 40 g
25			
30	<u>COLLECTOR 27</u> Reaction time: 115 min. 150°C Fumaric acid 116 g PEG 400 400 g Stearic acid 40 g		
35	In all synthesis other than 1 and 6, 1 ml concentrated sulphuric acid solved in 2 ml water was used as catalyst. In synthesis 1 no catalyst was used and in synthesis 6 methane sulphone acid was used. In table 1 below synthesis data is shown.		

TABLE 1

5	COLLECTOR	Blockage acid	SYNTHESIS TEMP. °C	Acid ratio	VISCOOSITY, mpas.23°C	CATALYST, T E O R . M W g/mole, synth. time. Min.
10	1	Tall oil	150	49.5	653	no,-,240
15	2	"	"	15.5	3210	yes,-,210
	3	"	175	8.1	5380	yes,-,165
	4	"	150	7.8	2640	yes,-,210
20	5	"	175	9.5	4960	yes,-,150
	6	"	200	11.3	2100	yes,-,180
25	7	"	"	7.7	>10000	yes,-,150
	8	"	150	15.1	>10000	yes,7000,210
	9	"	"	22.8	8720	yes,7000,195
30	10	"	175	12.5	>10000	yes,7000,155
	11	"	"	16.4	>10000	yes,7000,180
35	12	"	200	12.4	>10000	yes,7000,180
	13	"	"	12.8	>10000	yes,7000,180
	14	C-18	175	4.7	>10000	yes,-,180
40	15	"	150	7.6	>10000	yes,-,180
	16	"	200	3.1	>10000	yes,-,150
45	17	"	150	6.1	>10000	yes,-,180
	18	Tall oil	"	38.2	3710	yes,3000,210
	19	C-18	"	33.0	>10000	yes,3000,180
50	20	Tall oil	"	27.3	6830	yes,5000,210
	21	C-18	"	32.3	>10000	yes,5000,180
55	22	C-14	"	22.4	>10000	yes,7000,180
	23	C-12	"	31.5	>10000	yes,7000,180
	24	"	"	33.1	>10000	yes,3000,180
60	25	C-14	150	25.3	>10000	yes,7000,240
	26	C-18	200	5.4	>10000	yes,-,150
65	27	C-18	150	58	>10000	yes,7000,115

70 C-18 in the table signifies technical stearic acid
C-14 signifies myristic acid
C-12 signifies lauric acid

Polymer calculations

The amount of the included reactant was calculated through following method:

5 The average polymerisation ratio, X_n , which is the number of bifunctional molecules at the beginning of the reaction divided by the total number of polymer molecules, is calculated according to following:

10
$$X_n = (1+r)/(1+r-2xrxp) \quad (1)$$

but also according to

15
$$X_n = M_n/m \quad (2)$$

where M_n = the average molecular weight
 m = the molecular weight for the repeating unit in the molecule

20
$$r = N_A/N_B < 1$$

 p = ratio of transformation

If p is set to 1, then it is obtained:

25
$$X_n = (1+r)/(1-r) \implies r = (X_n - 1)(X_n + 1) \quad (3)$$

The amount of blockage acid is calculated according to following:

$$r = (N_A)/(N_A + 2N_B)$$

30 By stoic measurement balance $N_A = N_B$ and thereby becomes:

$$N_B = (N_B - N_B xr)/r \quad (4)$$

where N_A and N_B is the number of bifunctional molecules and 35 N_B is the number of monofunctional molecules, i.e. the blockage acid.

To calculate the number of blockage acid the desired molecular weight was firstly decided. Subsequently X_n

according to (2) was calculated. By means of (3) and (4) the number of blockage acid could be calculated later.

Acid ratio determination

5 Approximately 2g sample was weighed in a 300 ml E-flask. 100 ml ethanol was poured in a 300 ml E-flask and 15-20 drops phenolphthalein solution (0.05%) was added. The ethanol was titrated with 0.1M KOH until light rose alteration (only some drops) and was added to the saturated 10 test. Agitation was applied until the test was solved. The test solution was titrated until rose-coloured alteration and amount of required ml KOH was read off. The titration must be performed fast, since CO₂ from air is solved in the solution and uses KOH.

15

Calculation:

$$\text{Acid ratio} = M \times k \times V / m = 56.1 \times 0.1 \times V / m = 5.61 \times V / m$$

20 M = mole weight of potassiumhydroxide (56.1)
k = concentration of the KOH-solution
V = number of ml KOH consumed by the titration
m = weighted amount of sample in g

25 **Flotation experiment with the collector**

Daily and magazine newspapers were torn to pieces in approximately 4x4 cm pieces. After control of the dryness the newspapers and water as well as chemicals were collected in a pulper vessel. After slushing the pulp slurry is 30 allowed to rest after, which pH was controlled. Subsequently the slurry was further grinded with an Ultra-Turrax to obtain a more complete fibre release. The pulp was poured in the flotation cell and water was added. If the collector was not added in connection with the initial slushing we 35 added it before the flotation. The flotation was started. During the above mentioned steps we held a slurry temperature of 40 or 45°C, depending on flotations receipt.

pH was measured after the slushing. Sample for manufacturing brightness sheet and control of white water purity (see below) was taken before and after the flotation. Also, the sample was taken of the foam from the scrape.

5

At the tests 150 g paper raw material in 70% newspaper and 30% magazine paper, 2.5 l water (40°C), 3.2 g water-glass (46.8%) - 1% dosage of dry rich pulp, 1.5 g NaOH - 1% dosage, 0.45 g surfactant - 0.3% dosage (was reduced later to 0.075 g, i.e. 0.05% dosage) were used. The mixture diluted to 5% pulp concentration and slushed in a pulper for 5 min. It swelled subsequently in 1h after, which slushing occurred for 5 min. pH was measured until the mixture was diluted to 10 litre and stirred.

15

The slurry was poured in the flotation cell, which was filled with 40°C water. 1 litre was taken out for production of the brightness sheet and control of the white water purity. The flotation chemicals were added in form of 0.75 g collector - 0.5% dosage. At flotation with soap based collector 0.3 g - 0.2% dosage was added. Also, 1.78 g CaCl₂ was added to obtain 7°dH.

25 The flotation was accomplished in 30 min. After the flotation, 10 litres were tapped from the cell and the sample was taken for brightness sheet and the water phase.

Brightness sheet

Brightness sheet was made according to SCAN standard C11:75. 30 1 litre sample was taken out from the flotation cell and poured in mixer so that the fibres were separated from each other. The mixture diluted to 2 litre and pH adjusted to 4.0 - 5.5. 4 samples of 0.5 litres were sucked out in Büchner funnel with the diameter of 11 cm. The sheet was pressed or 35 rolled and dried in room temperature. The brightness was decided subsequently.

White water purity

The white water was controlled by following method: 200 ml suspension was taken out from the cell and fibres were filtered by a net basket. 60 ml of the filtrate was filtered 5 through Munktell filter paper type 00A (diameter = 5.5 cm). After drying was measured the brightness.

In table 2 the acid ratio as well as the brightness increase (Δ -brightness) for the different collectors are indicated.

10

TABLE 2 **Δ -brightness (ISO R457)**

	COLLECTOR	White water	Brightness sheet
15	1	-	2.11
20	2	-	2.12
	2	2.97	3.40
	3	-	2.85
25	4	3.17	2.48
	4	3.30	3.61
30	5	6.49	1.97
	6	4.33	3.90
	6	3.17	3.65
35	7	0.44	3.09
	7	3.46	3.17
40	8	2.38	3.63
	8	3.74	2.85
	9	4.24	3.66
45	10	7.77	4.70
	10	3.84	4.32
50	11	5.56	3.89
	0-sample	2.64	3.71
	0-sample	1.52	0.73
55	soap (0.2%)	5.57	5.14
	Soap (0.2%)	4.02	1.34
60	Tenneco (0.5%)	4.40	2.32
	Tenneco (0.5%)	2.05	1.90
	Bimex 400 (0.5%)	3.74	4.11

Result

Totally, 27 collectors were manufactured. By the attempts carried out, it was reviled that a collector based on 5 polyethylene glycol, maleine acid and stearic acid gave the best result. The molecular weight for the collector has been determine to approximately 6500 through HPLC. The clouding point was 57°C. The composition of the products and average 10 molecular weight was controlled by means of NMR, FTIR and HPLC.

The viscosity measurements point out that the molecular weight increases with the synthetic temperature. Possibly a certain cross linking at higher temperature. Small 15 differences in acid ratio but great differences in viscosity between the collector made at different synthetic temperatures indicates this.

The clouding point of the collector should be in few degrees 20 above the flotation temperature. We recommend that a number of different collectors are produced for optimal performance when the flotation temperature varies between different deinking plants.

25 It has not been able to indicate any connections between the flotation effect and the acid ratio. We have been able to discern better flotation result for products with higher viscosities. By comparisons between the different blockage acids the stearic acid has given best flotation result. Tall 30 oil as blockage acid has given uneven flotation result, which perhaps can be explained through its any complicated composition.

For the best collector, no 16, we optimized the dosage. Experiment with increasing concentration show that 0.3% 35 concentration collector gave best result. Higher amount gave poor flotation result. The results are shown in FIGURE 1. Comparing the experiment with a calcium stearate based

commercial collector, Tenneco[®], and the collector 16 according to the invention showed that, to attain same brightness, 4.5 units (ISO-R457), on brightness sheet as for a concentration of 0.35% collector 16, a concentration of 5 1.2% of Tenneco[®] was demanded, i.e. 3.5 times higher dosage. These results are shown in Fig. 2.

CLAIM

1. Collector intended to be used at deinking of waste paper according to the flotation process,
5 characterized therein,
that it includes a polyester obtained through reaction between 1) polyalkylene glycol, 2) di and/or tricarboxylic acid and/or anhydrides thereof as well as 3) a saturated fatty acid with 12-18 carbon atoms, and, which polyester
10 indicates a molecular weight between 3000-10000.
2. A collector according to claim 1,
characterized therein,
that the included source material shows merely aliphatic
15 carboxyl acids and alcohols.
3. A collector according to claim 2,
characterized therein,
that polyalkylene glycolen consists of polyethylene glycol,
20 polypropylene glycol or amalgamation thereof.
4. A collector according to any or some of proceeding claims,
characterized therein,
25 that di and tricarboxylic acid is preferably elected from the group consisting of maleine acid, fumaric acid, adipine acid, citric acid, oxalic acid and sebacic acid or anhydrides of these.
- 30 5. A collector according to any or some of proceeding claims,
characterized therein,
that it is produced through reaction between polyethylene glycol, maleine acid and stearic acid.

6. Use of a polyester obtained through reaction between 1) polyalkylene glycol, 2) di and/or tricarboxylic acid and/or anhydrides thereof as well as 3) saturated fatty acid with 12-18 carbon atoms, and, which polyester shows a molecular weight between 3000 - 10000, as collector for deinking waste paper according to the flotation process.

FIG. 1

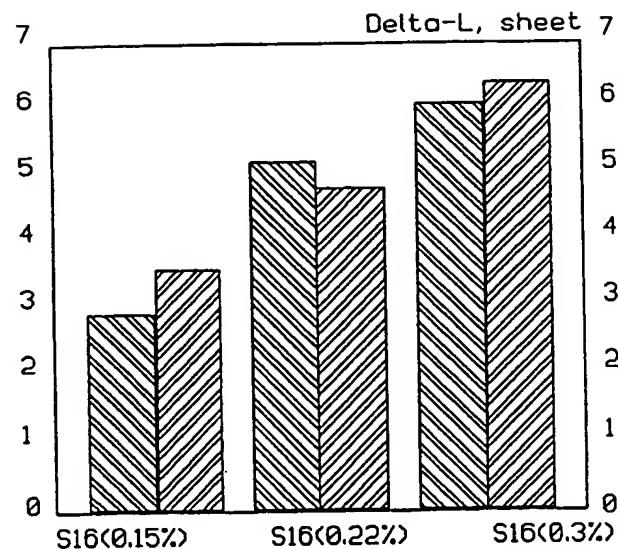
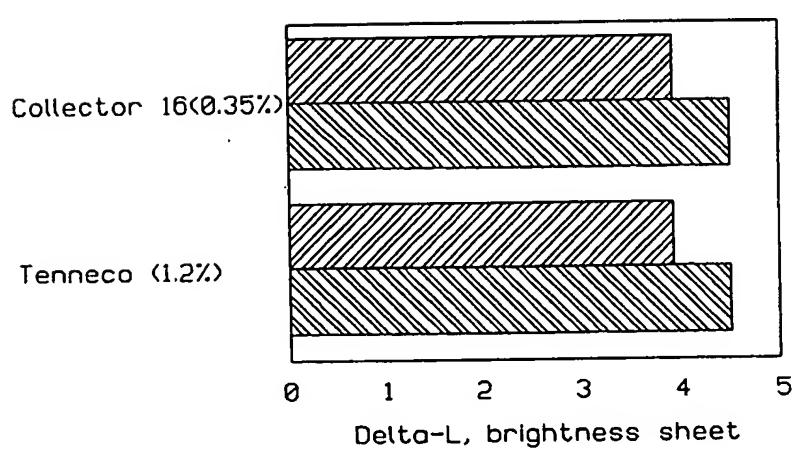


FIG. 2



INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 94/00547

A. CLASSIFICATION OF SUBJECT MATTER

IPC5: D21C 5/02

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC5: D21C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO, A1, 9103599 (BIM KEMI AB), 21 March 1991 (21.03.91) --	1-6
A	Dialog Information Services, File 351, DERWENT WPI, Dialog accession no.003525055, WPI accession no. 82-73041E/35, HONSHU PAPER MFG KK et al: "Removing ink from waste printed paper in flotation process using cpd. obtd. by addn. polymerising alkylene oxide with mono or polyfunctional alcohol and reac- ting the prod. with dicarboxylic acid; polyester pulp", JP 57117690, A, 820722, 8235 (Basic) -- -----	1-6

 Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier document but published on or after the international filing date	"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

Date of mailing of the international search report

6 October 1994

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INTERNATIONAL SEARCH REPORT
Information on patent family members

27/08/94

International application No.
PCT/SE 94/00547

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
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